

TITLE OF THE INVENTION

WAFER PROCESSING APPARATUS, WAFER PROCESSING METHOD,
AND SEMICONDUCTOR SUBSTRATE FABRICATION METHOD

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BACKGROUND OF THE INVENTION

FIELD OF THE INVENTION

The present invention relates to a wafer processing
apparatus, a wafer processing method, and a
10 semiconductor substrate fabrication method and, more
particularly, to a wafer processing apparatus for
processing a wafer by dipping it into a processing
solution, a wafer processing method, and a semiconductor
substrate fabrication method.

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DESCRIPTION OF THE RELATED ART

Cleaning processing is a typical example of wafer
processing. One subject of wafer cleaning is to increase
the speed. Japanese Patent Laid-Open No. 8-293478 has
20 disclosed a wafer cleaning method capable of increasing
the cleaning efficiency by supplying ultrasonic waves
while rotating a wafer, and an apparatus for practicing
this method.

The wafer cleaning method disclosed in Japanese
25 Patent Laid-Open No. 8-293478 is based on the

present invention is a wafer processing apparatus for processing a wafer by dipping the wafer into a processing solution, characterized by comprising a processing bath having a depth that allows to completely
5 dip the wafer into the processing solution, wafer rotating means for rotating one or a plurality of wafers held by a wafer holder by using a wafer rotating member which rotates about a shaft shifted from a portion immediately below a barycenter of the one or plurality
10 of wafers, and ultrasonic generating means for generating ultrasonic waves in the processing bath.

In the wafer processing apparatus, only the wafer rotating member is preferably arranged as a member for transmitting a rotating force to the wafer below the one
15 or plurality of wafers held by the wafer holder.

In the wafer processing apparatus, the wafer rotating member preferably comprises at least one rod member substantially parallel to the shaft, and the rod member preferably rotates about the shaft.

20 In the wafer processing apparatus, the rod member preferably has a diameter much smaller than a diameter of a cylinder virtually formed upon rotation of the rod member about the shaft.

In the wafer processing apparatus, the rod member
25 preferably has a groove which engages with a peripheral

portion of the wafer.

In the wafer processing apparatus, a section of the rod member taken along the shaft preferably has a substantially sine-wave shape.

5 In the wafer processing apparatus, a section of the rod member taken along the shaft preferably has a substantially full-wave rectifying shape.

In the wafer processing apparatus, the wafer rotating means preferably further comprises driving
10 force generating means arranged outside the processing bath, and driving force transmission means for transmitting a driving force generated by the driving force generating means to the wafer rotating member and rotating the wafer rotating member.

15 The wafer processing apparatus preferably further comprises a dividing member for dividing an interior of the processing bath into a processing wafer side and a side of the driving force transmission means.

In the wafer processing apparatus, the driving
20 force transmission means preferably transmits the driving force generated by the driving force generating means through a crank mechanism.

In the wafer processing apparatus, the processing bath preferably comprises a circulating mechanism having
25 an overflow bath.

In the wafer processing apparatus, the circulating mechanism preferably comprises contamination reducing means for reducing contamination of the wafer by particles.

5 In the wafer processing apparatus, the contamination reducing means preferably comprises a filter.

In the wafer processing apparatus, the contamination reducing means preferably comprises means
10 for adjusting flow of the processing solution in the processing bath.

In the wafer processing apparatus, the ultrasonic generating means preferably comprises an ultrasonic bath and an ultrasonic source, and the processing bath
15 preferably receives ultrasonic waves through an ultrasonic transmitting medium set in the ultrasonic bath.

The wafer processing apparatus preferably further comprises driving means for changing a relative
20 positional relationship between the ultrasonic source and a wafer to be processed.

In the wafer processing apparatus, the driving means preferably moves the ultrasonic source within the ultrasonic bath.

25 In the wafer processing apparatus, at least

A wafer processing method according to the present invention is a wafer processing method of processing a wafer while ultrasonic waves are supplied, characterized by comprising processing the wafer while entirely dipping the wafer into a processing solution and changing a position of an ultrasonic source.

The wafer processing method according to the present invention is characterized in that the wafer is cleaned using a wafer cleaning solution as the processing solution.

The wafer processing method is suitable for a method of etching the wafer using a wafer etching solution as the processing solution.

The wafer processing method is suitable for a
15 method of etching a porous silicon layer of a wafer
having the porous silicon layer using a porous silicon
etching solution as the processing solution.

The wafer processing method is suitable for a method of etching a porous silicon layer of a wafer having the porous silicon layer using, as the processing solution, any one of

(a) hydrofluoric acid,

(b) solution mixture prepared by adding at least one of alcohol and hydrogen peroxide to hydrofluoric acid,

[illegible]

comprising the step of forming a non porous layer on a porous layer formed on a surface of a first substrate, the step of bonding a first substrate side of a prospective structure and a second substrate prepared separately to sandwich the non porous layer between the first substrate side and the second substrate, the removal step of removing the first substrate from the bonded structure to expose the porous layer on a second substrate side thereof, and the etching step of etching the porous layer while the second substrate side on which the porous layer is exposed is completely dipped into an etching solution, and ultrasonic waves are supplied, thereby exposing surface of the second substrate side, the etching step rotating and vertically moving the second substrate side.

A semiconductor substrate fabrication method according to the present invention is characterized by comprising the step of forming a non porous layer on a porous layer formed on a surface of a first substrate, the step of bonding a first substrate side of a prospective structure and a second substrate prepared separately to sandwich the non porous layer between the first substrate side and the second substrate, the removal step of removing the first substrate from the bonded structure to expose the porous layer on a second

substrate side thereof, and the etching step of etching
the porous layer while the second substrate side on
which the porous layer is exposed is completely dipped
into an etching solution, and ultrasonic waves are
5 supplied, thereby exposing surface of the second
substrate side, the etching step changing a position of
an ultrasonic bath.

The etching solution used in the etching step is
preferably any one of

- 10 (a) hydrofluoric acid,
- (b) solution mixture prepared by adding at least
one of alcohol and hydrogen peroxide to hydrofluoric
acid,
- (c) buffered hydrofluoric acid,
- 15 (d) solution mixture prepared by adding at least
one of alcohol and hydrogen peroxide to buffered
hydrofluoric acid, and
- (e) solution mixture of hydrofluoric acid, nitric
acid, and acetic acid.

20 The removal step preferably comprises exposing the
porous layer by grinding, polishing, or etching the
first substrate from a back surface.

The removal step preferably comprises separating
the first substrate side and the second substrate side
25 at a boundary of the porous layer.

apparatus shown in Fig. 1;

Fig. 3 is a perspective view showing an example of the construction of a wafer rotating member;

Figs. 4A and 4B are views, respectively, showing the movement of a wafer when the wafer rotating member is rotated in a lifting direction;

Figs. 5A and 5B are views, respectively, showing the movement of a wafer having an orientation flat;

Figs. 6A and 6B are sectional views, respectively, showing another example of the construction of a wafer rotating rod;

Figs. 7A and 7B are sectional views, respectively, showing still another example of the construction of the wafer rotating rod;

Figs. 8A to 8C are views each showing an example of the shape of the section of the wafer rotating rod;

Fig. 9 is a view showing a mechanism for transmitting a driving torque generated by a motor to the rotating shaft of the wafer rotating member; and

Figs. 10A to 10F are views, respectively, showing the method of fabricating a semiconductor wafer.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Preferred embodiments of the present invention will be described in detail below with reference to the

accompanying drawings.

[First Embodiment]

Fig. 1 is a perspective view showing the schematic construction of a wafer processing apparatus according to the first embodiment of the present invention. Fig. 2 is a sectional view of the wafer processing apparatus shown in Fig. 1.

In a wafer processing apparatus 100 according to this embodiment, portions which may come into contact with a processing solution are preferably made from quartz or plastic in accordance with the intended use. Preferable examples of the plastic are a fluorine resin, vinyl chloride, polyethylene, polypropylene, polybutyleneterephthalate (PBT), and polyetheretherketone (PEEK). Preferable examples of the fluorine resin are PVDF, PFA, and PTFE.

This wafer processing apparatus 100 has a wafer processing bath 10, an overflow bath 20, an ultrasonic bath 30, and a wafer rotating mechanism (52 to 59) for rotating wafers 40.

To process wafers, the wafer processing bath 10 is filled with a processing solution (e.g., an etching solution or a cleaning solution). The overflow bath 20 for temporarily storing any processing solution overflowing from the wafer processing bath 10 is

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provided around the upper portion of the wafer
processing bath 10. The processing solution temporarily
stored in the overflow bath 20 is discharged from the
bottom portion of the overflow bath 20 to a circulator
5 21 through a discharge pipe 21a. The circulator 21
removes particles by filtering the discharged processing
solution and supplies the processing solution to the
bottom portion of the wafer processing bath 10 through a
supply pipe 21b. Consequently, particles in the wafer
10 processing bath 10 are efficiently removed.

The wafer processing bath 10 must have a depth by
which the wafers 40 are completely dipped. This prevents
particles from attaching to the wafers 40 at the
interface between the processing solution and ambient
15 atmosphere, and makes processing for the wafers 40
uniform.

When wafers are processed by completely dipping
them into the processing solution, and particles attach
to the wafers in the processing solution, the particles
20 easily return into the processing solution. However, if
only parts of wafers are dipped into the processing
solution, particles attaching to the wafers at the
interface between the processing solution and ambient
atmosphere are hardly removed from the wafers, and
25 exposed to ambient atmosphere while attaching to the

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rods 53 arranged substantially parallel to each other are coupled through connecting rods 54, and a rotating shaft 52 is coupled to almost the center of one connecting rod 54. The wafer rotating member 50 is
5 pivotally supported at the rotating shaft 52 by a shaft support portion 11. Note that another rotating shaft may be arranged on the side opposite to the rotating shaft 52.

The diameter of the wafer rotating rod 53 is set
10 much smaller than the diameter of a cylinder virtually formed upon rotation of the wafer rotating rods 53. With this setting, the transmission efficiency of a rotating torque and ultrasonic waves to the wafers 40 can be increased.

15 Standing waves, i.e., high- and low-strength portions of ultrasonic waves are usually formed between the bottom surface of the wafer processing bath 10 and the liquid surface. In this wafer processing apparatus 100, however, processing for the wafers 40 can be made
20 uniform because the wafers 40 are rotated while being vertically moved by rotation of the wafer rotating member 50.

Since the wafer rotating member 50 has the minimum member which interrupts the transmission of ultrasonic
25 waves between the bottom surface of the wafer processing

and can pinch the wafers 40. Therefore, a slip between the wafers 40 and the wafer rotating rod 53 is more effectively suppressed upon applying ultrasonic waves.

Further, since this wafer rotating rod 53 does not have any acute-angled portion, unlike the wafer rotating rod 53 shown in Figs. 6A and 6B, particles produced upon contact with the wafers 40 can be reduced. This effect can also be achieved by forming grooves 53c with a full-wave rectifying shape.

Figs. 8A, 8B, and 8C are views each showing an example of the shape of the section of the wafer rotating rod 53. The section of the wafer rotating rod 53 can have various shapes. For example, its section may have a circular shape as shown in Fig. 8A, an elliptic shape as shown in Fig. 8B, or a shape as shown in Fig. 8C.

The rotating shaft 52 of the wafer rotating member 50 is preferably shifted from a position immediately below the barycenter of the wafers 40 toward the side wall of the wafer holder 41 (x-axis direction).

Although the rotational direction of the wafer rotating rods 53 is not particularly limited, it is preferably a direction to lift the wafers 40 by the wafer rotating rod 53 closer to a position immediately below the barycenter of the wafers 40 (to be referred to

as the lifting direction hereinafter), as shown in
Fig. 2. This is because, if the wafer rotating rods 53
are rotated in the lifting direction, a force acts on
the wafers 40 substantially vertically, and hence
5 friction between the wafers 40 and the side wall of the
wafer holder 41 becomes small.

Figs. 4A and 4B are views, respectively, showing
the movement of the wafer 40 upon rotating the wafer
rotating member 50 in the lifting direction. A direction
10 A shows the lifting direction, and a direction B shows
the rotational direction of the wafer 40. The wafer 40
rotates in the direction B from the state in Fig. 4A
while being substantially vertically lifted by the wafer
rotating rod 53 on a side immediately below the
15 barycenter of the wafer 40. The wafer 40 passes through
the state shown in Fig. 4B, and returns to the state
shown in Fig. 4A after the wafer rotating rods 53 rotate
through 180°. Accordingly, the wafer 40 rotates while
swinging vertically.

20 Since the wafer rotating member 50 rotates so as to
virtually form a cylinder by the two wafer rotating rods
53, it can properly transmit a rotating force to even a
wafer having an orientation flat. Figs. 5A and 5B are
views, respectively, showing the movement of a wafer 40
25 having an orientation flat.

Not to interrupt the transmission of ultrasonic waves while the wafer 40 is efficiently rotated and vertically moved, the number of wafer rotating rods 53 is preferably two, as described above. However, the number of wafer rotating rods 53 may be one. Also in this case, the wafer 40 can be rotated and vertically moved. As far as the interruption of the transmission of ultrasonic waves can be allowed, the number of wafer rotating rods 53 may be three or more (for example, they are cylindrically laid out).

Fig. 9 is a view showing a mechanism for transmitting a driving torque generated by a motor 59 to the rotating shaft 52 of the wafer rotating member 50. The driving torque generated by the motor 59 is transmitted to a crank 55 via a crank 58 and connecting rods 57. One end of the crank 55 is coupled to the rotating shaft 52 so as to fit thereon, whereas the other end is pivotally supported by a bearing 58. The rotating shaft 52 is pivotally supported by a bearing portion 11a formed in the shaft support portion 11, and rotates upon reception of the driving torque transmitted through the crank 55.

The wafer rotating mechanism is not limited to the above construction, and suffices only to rotate the rotating shaft 52. For example, a bevel gear, a belt, or

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are tightly bonded at room temperature so as to sandwich the SiO_2 layer 504 between them (Fig. 10D). This bonding may be strengthened by performing anodic bonding, pressurization, or heat treatment, as needed, or a
5 combination of them.

When a single-crystal Si layer is formed as the non porous layer 503, the first substrate is preferably bonded to the second substrate 505 after the SiO_2 layer 504 is formed on the surface of the single-crystal Si
10 layer by thermal oxidization or the like.

Preferable examples of the second substrate 505 are an Si substrate, a substrate having an SiO_2 layer formed on an Si substrate, a light-transmitting substrate such as a quartz substrate or the like, and a sapphire
15 substrate. The second substrate 505 suffices to have a flat surface to be bonded, and may be another type of substrate.

Fig. 10D shows the bonded state of the first and second substrates via the SiO_2 layer 504. The SiO_2 layer
20 504 need not be formed when the non porous layer 503 or the second substrate is not Si.

In bonding, a thin insulating plate may be inserted between the first and second substrates.

The first substrate is removed from the second
25 substrate at the boundary of the porous Si layer 502

(Fig. 10E). The removal method includes the first method (of discarding the first substrate) using grinding, polishing, etching, or the like, and the second method of separating the first and second substrates at the boundary of the porous layer 502. In the second method, the first substrate can be recycled by removing porous Si left on the separated first substrate, and planarizing the surface of the first substrate, as needed.

10 The porous Si layer 502 is selectively etched and removed (Fig. 10F). The wafer processing apparatus 100 is suitable for this etching. Since this wafer processing apparatus supplies ultrasonic waves while completely dipping a wafer (in this case, the wafer shown in Fig. 10E) into an etching solution and moving (e.g., rotating or vertically moving) it, the wafer is hardly contaminated by particles, and the etching is made uniform. According to this wafer processing apparatus, the etching time is shortened, and the etching selectivity between the non porous layer 503 and the porous layer 504 increases. The etching time is shortened because etching is promoted by ultrasonic waves, and the etching selectivity increases because the promotion of etching by ultrasonic waves is more remarkable on the porous layer 504 than on the non

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porous layer 503.

When the non porous layer 503 is single-crystal Si, the following etching solutions are suited in addition to a general etching solution for Si.

5 (a) hydrofluoric acid

(b) solution mixture prepared by adding at least one of alcohol and hydrogen peroxide to hydrofluoric acid

(c) buffered hydrofluoric acid

10 (d) solution mixture prepared by adding at least one of alcohol and hydrogen peroxide to buffered hydrofluoric acid

(e) solution mixture of hydrofluoric acid, nitric acid, and acetic acid

15 Using these etching solutions, the porous layer 502 can be selectively etched to leave the underlying non porous layer 503 (single-crystal Si). The porous layer 502 is readily selectively etched by these etching solutions because porous Si has an enormous surface area and hence etching of the porous Si progresses at a very
20 high speed in comparison with the non porous Si layer.

Fig. 10E schematically shows a semiconductor substrate obtained by the above fabrication method. According to this fabrication method, the flat non
25 porous layer 503 (e.g., single-crystal Si layer) is

Silicon wafers were set in the wafer processing bath 10 filled with a solution mixture of ammonia, hydrogen peroxide, and ultrapure water at about 80°C. While the wafers were rotated, ultrasonic waves of about 1 MHz were applied to clean the wafers. By this cleaning, 95% or more of particles were removed from the wafer surfaces. Also, this removal of particles was done uniformly on the wafer surface.

[Example 3]

10 This example pertains to etching of a silicon layer.

Silicon wafers were set in the wafer processing bath 10 filled with a solution mixture prepared by mixing hydrofluoric acid, nitric acid, and acetic acid at a ratio of 1 : 200 : 200. While the wafers were rotated, ultrasonic waves of about 0.5 MHz were applied to etch the wafer surfaces for 30 sec. Consequently, the silicon wafers were uniformly etched by about 1.0 μm . The uniformity of the etching rate was $\pm 5\%$ or less on the wafer surface and between the wafers.

20 [Example 4]

This example relates to etching of an SiO_2 layer. Hydrofluoric acid is suitable for the etching of an SiO_2 layer.

Wafers on which an SiO_2 layer was formed were set in the wafer processing bath 10 filled with 1.2%

mixture of hydrofluoric acid, hydrogen peroxide, and ultrapure water. While the wafers were rotated, ultrasonic waves of about 0.25 MHz were applied to etch the porous silicon layer. Consequently, the porous silicon layer was uniformly etched by 5 μm . The uniformity of the etching rate was $\pm 3\%$ or less on the wafer surface and between the wafers.

Note that the mechanism of etching of porous silicon is disclosed in K. Sakaguchi et al., Jpn. J. Appl. Phys. Vol. 34, part 1, No. 2B, 842-847 (1995). According to this reference, porous silicon is etched when an etching solution penetrates into the pores of porous silicon by a capillary action and etches the walls of the pores. As the walls of the pores become thinner, these walls cannot support themselves beyond some point. Finally, the porous layer entirely collapses to complete the etching.

[Example 7]

This example concerns an SOI wafer fabrication method. Figs. 10A to 10F are sectional views showing the steps of the SOI wafer fabrication method according to this example.

First, a single-crystal Si substrate 501 for forming a first substrate was anodized in an HF solution to form a porous Si layer 502 (Fig. 10A). The

anodization conditions were as follows.

Current density : 7 (mA/cm²)

Anodizing solution : HF : H₂O : C₂H₅OH = 1 : 1 : 1

Time : 11 (min)

5 Porous Si thickness : 12 (μm)

Subsequently, the resultant substrate was allowed to oxidize in an oxygen atmosphere at 400°C for 1 h. By this oxidation, the inner walls of pores of the porous Si layer 502 were covered with a thermal oxide film.

10 A 0.30-μm thick single-crystal Si layer 503 was epitaxially grown on the porous Si layer 502 by a CVD (Chemical Vapor Deposition) process (Fig. 10B). The epitaxial growth conditions were as follows.

Source gas : SiH₂Cl₂/H₂

15 Gas flow rates : 0.5/180 (l/min)

Gas pressure : 80 (Torr)

Temperature : 950 (°C)

Growth rate : 0.3 (μm/min)

Next, a 200-nm thick SiO₂ layer 504 was formed on
20 the single-crystal Si layer (epitaxial layer) 503 by thermal oxidation (Fig. 10C).

The first substrate thus formed as shown in Fig. 10C and an Si substrate 505 as a second substrate were so bonded as to sandwich the SiO₂ layer 504
25 (Fig. 10D).

The single-crystal Si substrate 501 was removed from the first substrate to expose the porous Si layer 502 (Fig. 10E).

The wafers shown in Fig. 10E were set in the wafer processing bath 10 filled with a solution mixture of hydrofluoric acid, hydrogen peroxide, and ultrapure water. While the wafers were rotated, ultrasonic waves of about 0.25 MHz were applied to etch the porous Si layer 502 (Fig. 10F). The uniformity of the etching rate of the porous Si layer 502 was $\pm 5\%$ or less on the wafer surface and between the wafers. By applying ultrasonic waves while wafers are rotated as described above, it is possible to uniformly promote the collapse (etching) of porous Si on the wafer surface and between the wafers.

In the etching of the porous Si layer 502, the single-crystal Si layer (epitaxial layer) 503 functions as an etching stop layer. Therefore, the porous Si layer 502 is selectively etched on the entire surface of the wafer.

That is, the rate at which the single-crystal Si layer 503 is etched by the etching solution described above is very low, so the etching selectivity of the porous Si layer 502 to the single-crystal Si layer 503 is 10^5 or more. Accordingly, the etching amount of the single-crystal Si layer 503 is about a few tens of Å and

practically negligible.

Fig. 10F shows the SOI wafer obtained by the above steps. This SOI wafer has the 0.2- μm thick single-crystal Si layer 503 on the SiO_2 layer 504. The film thickness of this single-crystal Si layer 503 was measured at one hundred points over the entire surface and found to be $201 \text{ nm} \pm 4 \text{ nm}$.

In this example, a heat treatment was further performed in a hydrogen atmosphere at 1100°C for about 1 h. When the surface roughness of the resultant SOI wafers was evaluated with an atomic force microscope (AFM), the root-mean-square of the surface roughness in a square region of $5 \mu\text{m}$ side was about 0.2 nm. This quality is equivalent to that of common Si wafers on the market.

Also, after the above heat treatment the cross-sections of the SOI wafers were observed with a transmission electron microscope. As a consequence, no new crystal defects were produced in the single-crystal Si layer 503, indicating that high crystallinity was maintained.

It is possible to form an SiO_2 film on the single-crystal Si film (epitaxial layer) 503 of the first substance as described above, on the surface of the second substrate 505, or on both. In any of these

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cases, results similar to these described above were obtained.

Furthermore, even when a light-transmitting wafer such as a quartz wafer was used as the second substrate, a high-quality SOI wafer could be formed by the above fabrication steps. However, the heat treatment in the hydrogen atmosphere was performed at a temperature of 1,000°C or less in order to prevent slip in the single-crystal Si layer 503 caused by the difference between the thermal expansion coefficients of the quartz (second substrate) and the single-crystal Si layer 503. [Example 8]

This example is directed to another SOI wafer fabrication method. Fabrication steps which can be expressed by drawings are the same as those shown in Figs. 10A to 10F, so the method will be described below with reference to Figs. 10A to 10F.

First, a single-crystal Si substrate 501 for forming a first substrate was anodized in an HF solution to form a porous Si layer 502 (Fig. 10A). The anodization conditions were as follows.

First stage:

Current density: 7 (mA/cm²)

Anodizing solution : HF : H₂O : C₂H₅OH = 1 : 1 : 1

Time : 5 (min)

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Porous Si thickness : 5.5 (μm)

Second stage:

Current density : 21 (mA/cm^2)

Anodizing solution : HF : H₂O : C₂H₅OH = 1 : 1 : 1

5 Time : 20 (sec)

Porous Si thickness : 0.5 (μm)

Subsequently, the resultant substrate was allowed to oxidize in an oxygen atmosphere at 400°C for 1 h. By this oxidation, the inner walls of pores of the porous
10 Si layer 502 were covered with a thermal oxide film.

A 0.15- μm thick single-crystal Si layer 503 was epitaxially grown on the porous Si layer 502 by a CVD (Chemical Vapor Deposition) process (Fig. 10B). The epitaxial growth conditions were as follows.

15 Source gas : SiH₂Cl₂/H₂

Gas flow rates : 0.5/180 (l/min)

Gas pressure : 80 (Torr)

Temperature : 950 (°C)

Growth rate : 0.3 ($\mu\text{m}/\text{min}$)

20 Next, a 100-nm thick SiO₂ layer 504 was formed on the single-crystal Si layer (epitaxial layer) 503 by oxidation (Fig. 10C).

The first substrate thus formed as shown in Fig. 10C and a second Si substrate 505 were so bonded as
25 to sandwich the SiO₂ layer 504 (Fig. 10D).

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The bonded wafers was separated into two wafers from the porous Si layer formed at a current density of 21 mA/cm² (second stage), thereby exposing the porous Si layer 503 to the surface of the second substrate 505 (Fig. 10E). Examples of the method of separating the bonded wafers are 1) mechanically pulling the two substrates, 2) twisting the substrates, 3) pressurizing the substrates, 4) driving a wedge between the substrates, 5) peeling the substrates by oxidizing from their end faces, 6) using thermal stress, and 7) applying ultrasonic waves, and it is possible to selectively use any of these methods.

The wafers shown in Fig. 10E were set in the wafer processing bath 10 filled with a solution mixture of hydrofluoric acid, hydrogen peroxide, and ultrapure water. While the wafers were rotated, ultrasonic waves of about 0.25 MHz were applied to etch the porous Si layer 502 (Fig. 10F). The uniformity of the etching rate of the porous Si layer 502 was $\pm 5\%$ or less on the wafer surface and between the wafers. By applying ultrasonic waves while wafers are rotated as described above, it is possible to uniformly promote the collapse (etching) of porous Si on the wafer surface and between the wafers.

In the etching of the porous Si layer 502, the single-crystal Si layer (epitaxial layer) 503 functions

as an etching stop layer. Therefore, the porous Si layer 502 is selectively etched on the entire surface of the wafer.

That is, the rate at which the single-crystal Si layer 503 is etched by the etching solution described above is very low, so the etching selectivity of the porous Si layer 502 to the single-crystal Si layer 503 is 10^5 or more. Accordingly, the etching amount of the single-crystal Si layer 503 is about a few tens of Å and practically negligible.

Fig. 10F shows the SOI wafer obtained by the above steps. This SOI wafer has the 0.1-μm thick single-crystal Si layer 503 on the SiO₂ layer 504. The film thickness of this single-crystal Si layer 503 was measured at one hundred points over the entire surface and found to be 101 nm ± 3 nm.

In this example, a heat treatment was further performed in a hydrogen atmosphere at 1,100°C for about 1 h. When the surface roughness of the resultant SOI wafers was evaluated with an atomic force microscope (AFM), the root-mean-square of the surface roughness in a square region of 5 μm side was about 0.2 nm. This quality is equivalent to that of common Si wafers on the market.

Also, after the above heat treatment the cross-

sections of the SOI wafers were observed with a transmission electron microscope. As a consequence, no new crystal defects were produced in the single-crystal Si layer 503, indicating that high crystallinity was maintained.

It is possible to form an SiO_2 film on the single-crystal Si film (epitaxial layer) 503 of the first substrate as described above, on the surface of the second substrate 505, or on both. In any of these cases, results similar to these described above were obtained.

Furthermore, even when a light-transmitting wafer such as a quartz wafer was used as the second substrate, a high-quality SOI wafer could be formed by the above fabrication steps. However, the heat treatment in the hydrogen atmosphere was performed at a temperature of $1,000^\circ\text{C}$ or less in order to prevent slip in the single-crystal Si layer 503 caused by the difference between the thermal expansion coefficients of the quartz (second substrate) and the single-crystal Si layer 503.

In this example, the first substrate (to be referred to as the separated substrate hereinafter) obtained by separating the bonded wafers into two wafers can be reused. That is, the separated substrate can be reused as the first or second substrate by selectively

etching the porous Si film remaining on the surface of
the substrate by the same etching method as for the
porous Si film described above and processing the
resultant material (e.g., annealing in a hydrogen
5 processing or a surface treatment such as surface
polishing).

In examples 7 and 8 described above, epitaxial
growth is used to form a single-crystal Si layer on a
porous Si layer. However, it is also possible to use
10 other various methods such as CVD, MBE, sputtering, and
liquid phase growth in the formation of a single-crystal
Si layer.

Also, a semiconductor layer of a single-crystal
compound such as GaAs or InP can be formed on a porous
15 Si layer by epitaxial growth. If this is the case,
wafers suited to high-frequency devices such as "GaAs on
Si" and "GaAs on Glass (Quartz)" and QEIC can be made.

Furthermore, although a solution mixture of 49%
hydrofluoric acid and 30% hydrogen peroxide is suitable
20 for an etching solution for selectively etching a porous
Si layer, the following etching solutions are also
suited. This is so because porous Si has an enormous
surface area and hence can be readily selectively etched.

- (a) hydrofluoric acid
- 25 (b) solution mixture prepared by adding at least

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one of alcohol and hydrogen peroxide to hydrofluoric acid

(c) buffered hydrofluoric acid

(d) solution mixture prepared by adding at least
5 one of alcohol and hydrogen peroxide to buffered hydrofluoric acid

(e) solution mixture of hydrofluoric acid, nitric acid, and acetic acid

Note that the other fabrication steps are not
10 limited to the conditions in the above examples, and so other various conditions can be used.

The present invention can reduce contamination of wafers by particles and make wafer processing uniform.

The present invention is not limited to the above
15 embodiments and various changes and modifications can be made within the spirit and scope of the present invention. Therefore, to apprise the public of the scope of the present invention the following claims are made.